Applicant: Michele Coutu Hresko et al. Attorney's Docket No.: 12557-015001

Serial No.: 10/771,708 Filed: February 4, 2004

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In the specification:

Insert the paper copy of the Sequence Listing filed herewith following the Drawings.

Replace the paragraph beginning at page 36, line 3 with the following rewritten paragraph:

C. elegans were grown on lawns of E. coli genetically engineered to produce double-stranded RNA (dsRNA) designed to inhibit PANZP1 or PANZP2 expression in order to investigate whether PANZP1 or PANZP2 expression is essential. Briefly, E. coli were transformed with genomic fragments encoding portions of the C. elegans PANZP1 or the PANZP2 gene. A 1048 nucleotide fragment was amplified from the PANZP1 gene using oligo-nucleotide primers containing the sequences 5'-TCAGTGACGTTATGTCCTCC-3' (SEQ ID NO: [21]51) and 5'-TGACAGATGGAACATTCTCC-3' (SEQ ID NO: [22]52). A 926 nucleotide fragment was amplified from the PANZP2 gene using oligo -nucleotide primers containing the sequences 5'-ACTTCAGGACACGACTTGAC-3' (SEQ ID NO: [23]53) and 5'-CAATCAGAGATGGTAACTCC-3' (SEQ ID NO: [24]54) respectively. The cloned PANZP1 and PANZP2 genomic fragments were cloned separately into an E. coli expression vector between opposing T7 polymerase promoters. The expression clones were separately transformed into a strain of E. coli that carries an IPTG-inducible T7 polymerase. As a control, E. coli was transformed with a gene encoding the Green Fluorescent Protein (GFP).